

10/823,211

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PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

* * * * * Welcome to STN International * * * * *

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NEWS	2	"Ask CAS" for self-help around the clock
NEWS	3 FEB 25	CA/CAPLUS - Russian Agency for Patents and Trademarks (ROSPATENT) added to list of core patent offices covered
NEWS	4 FEB 28	PATDPAFULL - New display fields provide for legal status data from INPADOC
NEWS	5 FEB 28	BABS - Current-awareness alerts (SDIs) available
NEWS	6 FEB 28	MEDLINE/LMEDLINE reloaded
NEWS	7 MAR 02	GBFULL: New full-text patent database on STN
NEWS	8 MAR 03	REGISTRY/ZREGISTRY - Sequence annotations enhanced
NEWS	9 MAR 03	MEDLINE file segment of TOXCENTER reloaded
NEWS	10 MAR 22	KOREAPAT now updated monthly; patent information enhanced
NEWS	11 MAR 22	Original IDE display format returns to REGISTRY/ZREGISTRY
NEWS	12 MAR 22	PATDPASPC - New patent database available
NEWS	13 MAR 22	REGISTRY/ZREGISTRY enhanced with experimental property tags
NEWS	14 APR 04	EPFULL enhanced with additional patent information and new fields
NEWS	15 APR 04	EMBASE - Database reloaded and enhanced
NEWS	16 APR 18	New CAS Information Use Policies available online
NEWS	17 APR 25	Patent searching, including current-awareness alerts (SDIs), based on application date in CA/CAPLUS and USPATFULL/USPAT2 may be affected by a change in filing date for U.S. applications.
NEWS	18 APR 28	Improved searching of U.S. Patent Classifications for U.S. patent records in CA/CAPLUS
NEWS EXPRESS	JANUARY 10	CURRENT WINDOWS VERSION IS V7.01a, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 10 JANUARY 2005
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NEWS INTER	General Internet Information	
NEWS LOGIN	Welcome Banner and News Items	
NEWS PHONE	Direct Dial and Telecommunication Network Access to STN	
NEWS WWW	CAS World Wide Web Site	(general information)

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* * * * * STN Columbus * * * * *

=> file req

SINCE FILE	TOTAL
ENTRY	SESSION
0.21	0.21

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

New CAS Information Use Policies, enter HELP USAGETERMS for details.

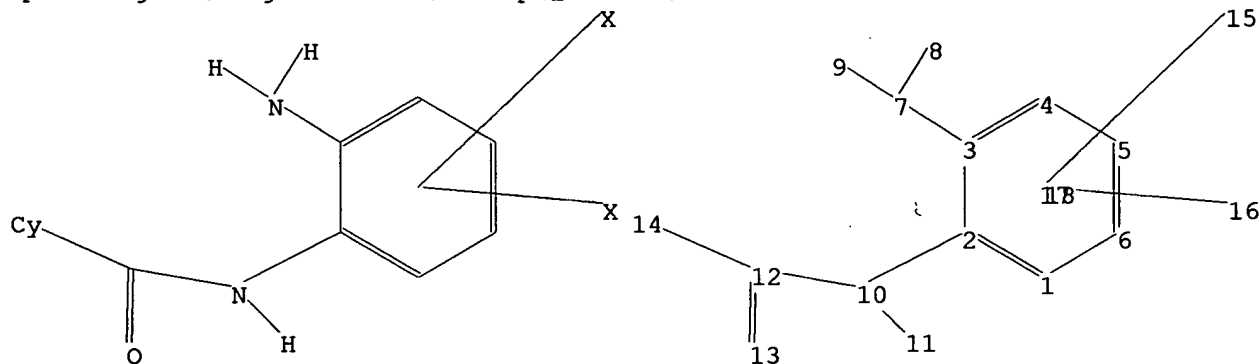
Please note that search-term pricing does apply when conducting SmartSELECT searches.

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*
* The CA roles and document type information have been removed from
* the IDE default display format and the ED field has been added,
* effective March 20, 2005.  A new display format, IDERL, is now
* available and contains the CA role and document type information.
*
*****
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Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at:
<http://www.cas.org/ONLINE/DBSS/registryss.html>

 \Rightarrow

Uploading C:\Program Files\Stnexp\Queries\10823211.str



chain nodes :

7 8 9 10 11 12 13 14 15 16

10/823,211

ring nodes :

1 2 3 4 5 6

chain bonds :

2-10 3-7 7-8 7-9 10-11 10-12 12-13 12-14

ring bonds :

1-2 1-6 2-3 3-4 4-5 5-6

exact/norm bonds :

2-10 3-7 10-12 12-13 12-14

exact bonds :

7-8 7-9 10-11

normalized bonds :

1-2 1-6 2-3 3-4 4-5 5-6

Match level :

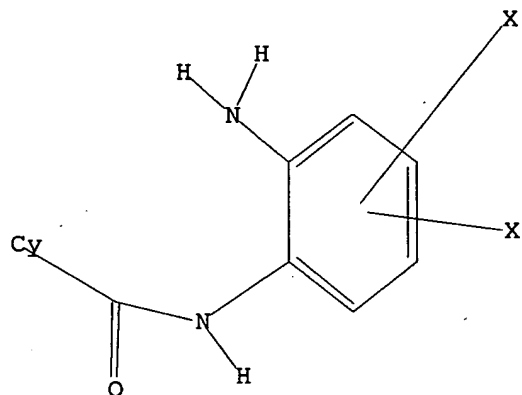
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:CLASS 8:CLASS 9:CLASS 10:CLASS
11:CLASS 12:CLASS 13:CLASS 14:Atom 15:CLASS 16:CLASS 17:CLASS 18:CLASS

L1 STRUCTURE UPLOADED

=> d l1

L1 HAS NO ANSWERS

L1 STR



Structure attributes must be viewed using STN Express query preparation.

=> s l1 sss sam

SAMPLE SEARCH INITIATED 13:11:11 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 919 TO ITERATE

100.0% PROCESSED 919 ITERATIONS

2 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 16562 TO 20198

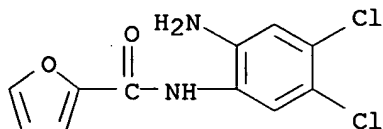
PROJECTED ANSWERS: 2 TO 124

L2 2 SEA SSS SAM L1

=> d scan

10/823,211

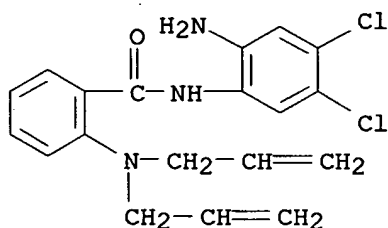
L2 2 ANSWERS REGISTRY COPYRIGHT 2005 ACS on STN
IN 2-Furancarboxamide, N-(2-amino-4,5-dichlorophenyl)- (9CI)
MF C11 H8 Cl2 N2 O2



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):1

L2 2 ANSWERS REGISTRY COPYRIGHT 2005 ACS on STN
IN Benzamide, N-(2-amino-4,5-dichlorophenyl)-2-(di-2-propenylamino)- (9CI)
MF C19 H19 Cl2 N3 O



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

ALL ANSWERS HAVE BEEN SCANNED

=> s l1 sss ful
FULL SEARCH INITIATED 13:11:35 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 19120 TO ITERATE

100.0% PROCESSED 19120 ITERATIONS
SEARCH TIME: 00.00.01

51 ANSWERS

L3 51 SEA SSS FUL L1

=> file caplus

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

161.76

161.97

FILE 'CAPLUS' ENTERED AT 13:11:41 ON 09 MAY 2005
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
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10/823,211

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FILE COVERS 1907 - 9 May 2005 VOL 142 ISS 20
FILE LAST UPDATED: 8 May 2005 (20050508/ED)

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

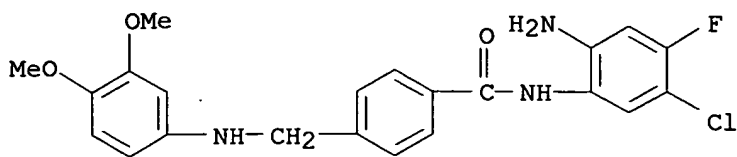
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L4 14 L3

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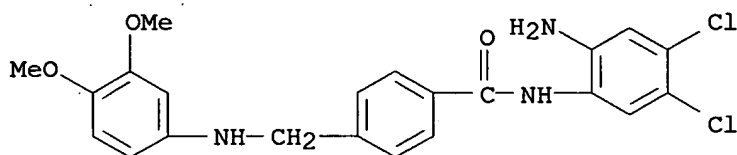
L4 ANSWER 1 OF 14 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2005:300395 CAPLUS
DN 142:355054
TI Preparation of amide derivatives as inhibitors of histone deacetylase
IN Moradei, Oscar; Paquin, Isabelle; Leit, Silvana; Frechette, Sylvie; Vaisburg, Arkadii; Besterman, Jeffrey M.; Tessier, Pierre; Mallais, Tammy C.
PA Methylgene, Inc., Can.
SO PCT Int. Appl., 559 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2005030705	A1	20050407	WO 2004-US31591	20040924
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
	RW:	BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
PRAI	US 2003-505884P	P	20030924		
	US 2003-532973P	P	20031229		
	US 2004-561082P	P	20040409		
IT	849235-31-0P 849235-32-1P				
	RL:	PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)			
		(preparation of amide derivs. as inhibitors of histone deacetylase)			
RN	849235-31-0 CAPLUS				
CN	Benzamide, N-(2-amino-5-chloro-4-fluorophenyl)-4-[[[3,4-dimethoxyphenyl]amino]methyl]- (9CI) (CA INDEX NAME)				



RN 849235-32-1 CAPLUS

CN Benzamide, N-(2-amino-4,5-dichlorophenyl)-4-[[[3,4-dimethoxyphenyl]amino]methyl]- (9CI) (CA INDEX NAME)



RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 14 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2005:300394 CAPLUS

DN 142:373563

TI Preparation of amide derivatives as inhibitors of histone deacetylase

IN Moradei, Oscar; Paquin, Isabelle; Leit, Silvana; Frechette, Sylvie;
Vaisburg, Arkadii; Besterman, Jeffrey M.; Tessier, Pierre; Mallais, Tammy
C.

PA Methylgene, Inc., Can.

SO PCT Int. Appl., 389 pp.
CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2005030704	A1	20050407	WO 2004-US31590	20040924
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	CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,				
	GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,				
	LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI,				
	NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY,				
	TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW:				
	BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM,				
	AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK,				
	EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE,				
	SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE,				
	SN, TD, TG				
PRAI	US 2003-505884P	P	20030924		
	US 2003-532973P	P	20031229		
	US 2004-561082P	P	20040409		

IT 849235-31-0P 849235-32-1P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU
(Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
(Uses)

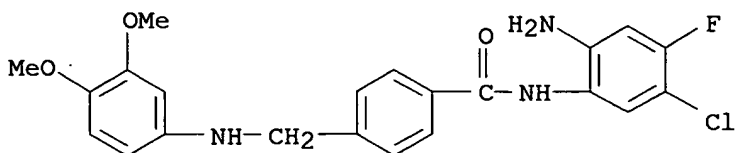
(preparation of amide derivs. as inhibitors of histone deacetylase)

RN 849235-31-0 CAPLUS

CN Benzamide, N-(2-amino-5-chloro-4-fluorophenyl)-4-[[[3,4-

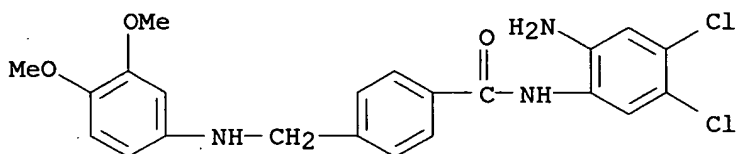
10/823,211

dimethoxyphenyl)amino]methyl]- (9CI) (CA INDEX NAME)



RN 849235-32-1 CAPLUS

CN Benzamide, N-(2-amino-4,5-dichlorophenyl)-4-[[3,4-dimethoxyphenyl]amino]methyl]- (9CI) (CA INDEX NAME)



RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 3 OF 14 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2004:654774 CAPLUS

DN 141:190788

TI A preparation of N-containing heterocyclic compounds, useful as vanilloid receptor ligands

IN Doherty, Elizabeth M.; Fotsch, Christopher H.; Han, Nianhe; Hungate, Randall W.; Liu, Qingyian; Norman, Mark H.; Xi, Ning; Xu, Shimin

PA Amgen Inc., USA

SO U.S. Pat. Appl. Publ., 38 pp.

CODEN: USXXCO

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2004157845	A1	20040812	US 2004-775980	20040209
	WO 2004072068	A1	20040826	WO 2004-US3908	20040209
	W:	AE, AE, AG, AL, AL, AM, AM, AM, AT, AT, AU, AZ, AZ, BA, BB, BG, BG, BR, BR, BW, BY, BY, BZ, BZ, CA, CH, CN, CN, CO, CO, CR, CR, CU, CU, CZ, CZ, DE, DE, DK, DK, DM, DZ, EC, EC, EE, EE, EG, ES, ES, FI, FI, GB, GD, GE, GE, GH, GM, HR, HR, HU, HU, ID, IL, IN, IS, JP, JP, KE, KE, KG, KG, KP, KP, KR, KR, KZ, KZ, KZ, LC, LK, LR, LS, LS, LT, LU, LV, MA, MD, MD, MG, MK, MN, MW, MX, MX, MZ, MZ, NA, NI			
	RW:	BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			

PRAI US 2003-446511P P 20030210

OS MARPAT 141:190788

IT 736971-85-0P

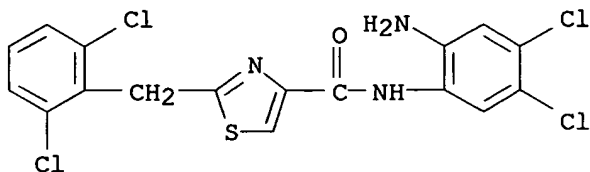
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of heterocyclic compds., useful as vanilloid receptor ligands)

RN 736971-85-0 CAPLUS

10/823,211

CN 4-Thiazolecarboxamide, N-(2-amino-4,5-dichlorophenyl)-2-[(2,6-dichlorophenyl)methyl]- (9CI) (CA INDEX NAME)



L4 ANSWER 4 OF 14 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2004:589250 CAPLUS

DN 141:140470

TI Preparation of aminophenylbenzamides as inhibitors of histone deacetylase

IN Delorme, Daniel; Zhou, Zhihong

PA Methylgene, Inc., Can.

SO U.S. Pat. Appl. Publ., 318 pp., Cont.-in-part of U.S. Ser. No. 242,304.

CODEN: USXXCO

DT Patent

LA English

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2004142953	A1	20040722	US 2003-358556	20030204
	US 2004106599	A1	20040603	US 2002-242304	20020912
	WO 2004069823	A1	20040819	WO 2004-CA139	20040204
	W:	AE, AE, AG, AL, AL, AM, AM, AM, AT, AT, AU, AZ, AZ, BA, BB, BG, BG, BR, BR, BW, BY, BY, BZ, BZ, CA, CH, CN, CN, CO, CO, CR, CR, CU, CU, CZ, CZ, DE, DE, DK, DK, DM, DZ, EC, EC, EE, EE, EG, ES, ES, FI, FI, GB, GD, GE, GE, GH, GM, HR, HR, HU, HU, ID, IL, IN, IS, JP, JP, KE, KE, KG, KG, KP, KP, KR, KR, KZ, KZ, LC, LK, LR, LS, LS, LT, LU, LV, MA, MD, MD, MG, MK, MN, MW, MX, MX, MZ, MZ, NA, NI			
	RW:	BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
PRAI	US 2001-322402P	P	20010914		
	US 2002-391728P	P	20020626		
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	US 2003-358556	A	20030204		

OS MARPAT 141:140470

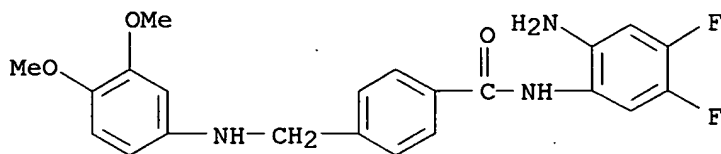
IT 503043-94-5P, N-(2-Amino-4,5-difluorophenyl)-4-[[(3,4-dimethoxyphenyl)amino]methyl]benzamide

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

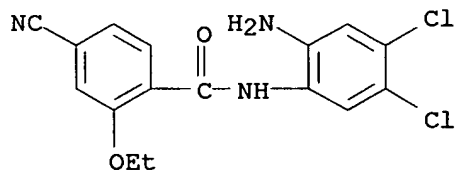
(drug candidate; preparation of aminophenylbenzamides as inhibitors of histone deacetylase for treating cell proliferative disorders)

RN 503043-94-5 CAPLUS

CN Benzamide, N-(2-amino-4,5-difluorophenyl)-4-[[(3,4-dimethoxyphenyl)amino]methyl]- (9CI) (CA INDEX NAME)



L4 ANSWER 5 OF 14 CAPLUS COPYRIGHT 2005 ACS on STN
 AN 2003:311219 CAPLUS
 DN 140:14257
 TI Synthesis of photoactivable inhibitors of osteoclast vacuolar ATPase
 AU Biasotti, Barbara; Dallavalle, Sabrina; Merlini, Lucio; Farina, Carlo;
 Gagliardi, Stefania; Parini, Carlo; Belfiore, Pietro
 CS Dipartimento di Scienze Molecolari Agroalimentari, Università di Milano,
 Milan, 20133, Italy
 SO Bioorganic & Medicinal Chemistry (2003), 11(10), 2247-2254
 CODEN: BMECEP; ISSN: 0968-0896
 PB Elsevier Science Ltd.
 DT Journal
 LA English
 IT **316810-13-6P**, N-(2-Amino-4,5-dichlorophenyl)-2-ethoxy-4-
 cyanobenzamide
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
 (Reactant or reagent)
 (preparation and osteoclast vacuolar ATPase inhibitory activity of
 photoactivable inhibitors containing trifluoroaziridinophenyl or
 benzophenone moiety and iodine)
 RN 316810-13-6 CAPLUS
 CN Benzamide, N-(2-amino-4,5-dichlorophenyl)-4-cyano-2-ethoxy- (9CI) (CA
 INDEX NAME)

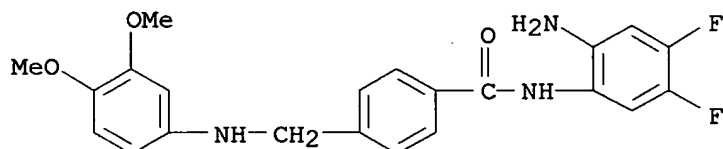


RE.CNT 24 THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 6 OF 14 CAPLUS COPYRIGHT 2005 ACS on STN
 AN 2003:242160 CAPLUS
 DN 138:271705
 TI Preparation of triazinyl and other carboxamides as inhibitors of histone
 deacetylase
 IN Delorme, Daniel; Woo, Soon Hyung; Vaisburg, Arkadii; Moradel, Oscar; Leit,
 Silvana; Raepfel, Stephane; Frechette, Sylvie; Bouchain, Giliane
 PA Methylgene, Inc., Can.
 SO PCT Int. Appl., 347 pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	WO 2003024448	A2	20030327	WO 2002-US29017	20020912

WO 2003024448 A3 20031113
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
EP 1429765 A2 20040623 EP 2002-763627 20020912
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK
BR 2002012510 A 20040824 BR 2002-12510 20020912
JP 2005508905 T2 20050407 JP 2003-528544 20020912
PRAI US 2001-322402P P 20010914
US 2002-391728P P 20020626
WO 2002-US29017 W 20020912
OS MARPAT 138:271705
IT 503043-94-5P, N-(2-Amino-4,5-difluorophenyl)-4-[[(3,4-dimethoxyphenyl)amino]methyl]benzamide
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of triazinyl and other carboxamides as inhibitors of histone deacetylase for treating cell proliferative disorders)
RN 503043-94-5 CAPLUS
CN Benzamide, N-(2-amino-4,5-difluorophenyl)-4-[[(3,4-dimethoxyphenyl)amino]methyl]- (9CI) (CA INDEX NAME)



L4 ANSWER 7 OF 14 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:133003 CAPLUS
DN 138:175871
TI N-monoacylated derivatives of o-phenylenediamines and their six-membered heterocyclic analogs as antitumor agents
IN Haag, Rainer; Leser-Reiff, Ulrike; Limberg, Anja; Weidner, Michael; Zimmermann, Gerd
PA F. Hoffmann-La Roche AG, Switz.
SO PCT Int. Appl., 27 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003013484	A2	20030220	WO 2002-EP8708	20020805
	WO 2003013484	A3	20030417		
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,			

LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZM, ZW

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

EP 1416928 A2 20040512 EP 2002-767319 20020805

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK

JP 2005508311 T2 20050331 JP 2003-518494 20020805

US 2003139404 A1 20030724 US 2002-212901 20020806

US 6869953 B2 20050322

US 2004192744 A1 20040930 US 2004-823211 20040413

PRAI EP 2001-118741 A 20010807

WO 2002-EP8708 W 20020805

US 2002-212901 A1 20020806

OS MARPAT 138:175871

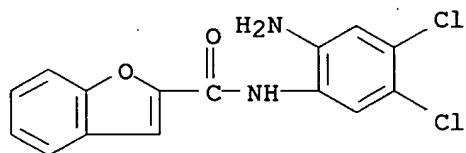
IT 497825-42-0P 497825-43-1P 497825-44-2P
 497825-45-3P 497825-46-4P 497825-47-5P
 497825-48-6P 497825-49-7P 497825-50-0P
 497825-51-1P 497825-52-2P 497825-53-3P
 497825-54-4P 497825-55-5P 497825-56-6P
 497825-57-7P 497825-58-8P 497825-59-9P
 497825-60-2P 497825-61-3P 497825-62-4P
 497825-63-5P 497825-64-6P 497825-65-7P
 497825-66-8P 497825-67-9P 497825-68-0P
 497825-69-1P 497825-70-4P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of monoacylated derivs. of o-phenylenediamines and their 6-membered heterocyclic analogs as antitumor agents)

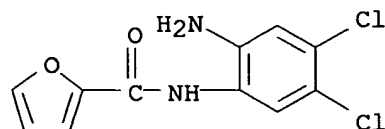
RN 497825-42-0 CAPLUS

CN 2-Benzofurancarboxamide, N-(2-amino-4,5-dichlorophenyl)- (9CI) (CA INDEX NAME)



RN 497825-43-1 CAPLUS

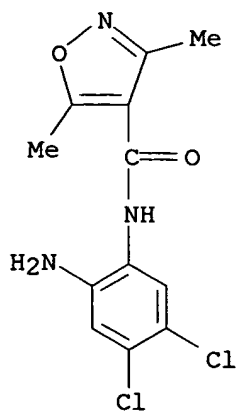
CN 2-Furancarboxamide, N-(2-amino-4,5-dichlorophenyl)- (9CI) (CA INDEX NAME)



RN 497825-44-2 CAPLUS

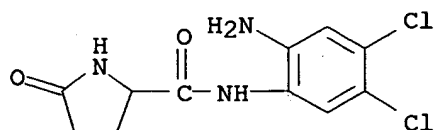
CN 4-Isioxazolecarboxamide, N-(2-amino-4,5-dichlorophenyl)-3,5-dimethyl- (9CI) (CA INDEX NAME)

10/823,211



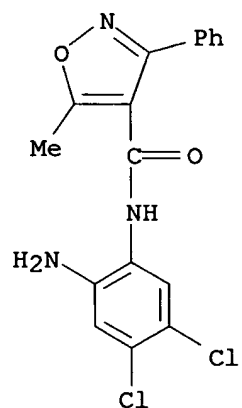
RN 497825-45-3 CAPLUS

CN 2-Pyrrolidinecarboxamide, N-(2-amino-4,5-dichlorophenyl)-5-oxo- (9CI) (CA INDEX NAME)



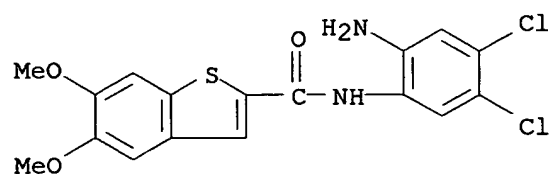
RN 497825-46-4 CAPLUS

CN 4-Isoxazolecarboxamide, N-(2-amino-4,5-dichlorophenyl)-5-methyl-3-phenyl- (9CI) (CA INDEX NAME)



RN 497825-47-5 CAPLUS

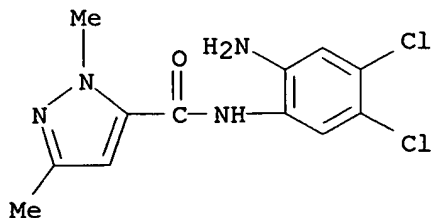
CN Benzo[b]thiophene-2-carboxamide, N-(2-amino-4,5-dichlorophenyl)-5,6-dimethoxy- (9CI) (CA INDEX NAME)



10/823,211

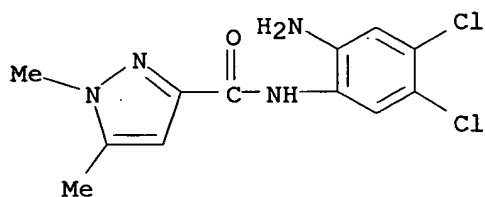
RN 497825-48-6 CAPLUS

CN 1H-Pyrazole-5-carboxamide, N-(2-amino-4,5-dichlorophenyl)-1,3-dimethyl-
(9CI) (CA INDEX NAME)



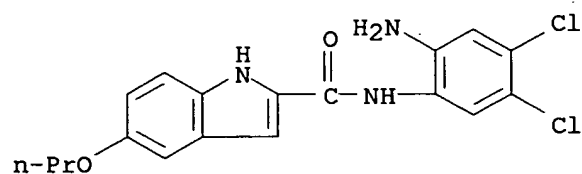
RN 497825-49-7 CAPLUS

CN 1H-Pyrazole-3-carboxamide, N-(2-amino-4,5-dichlorophenyl)-1,5-dimethyl-
(9CI) (CA INDEX NAME)



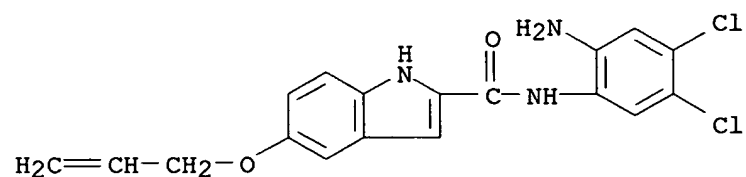
RN 497825-50-0 CAPLUS

CN 1H-Indole-2-carboxamide, N-(2-amino-4,5-dichlorophenyl)-5-propoxy- (9CI)
(CA INDEX NAME)



RN 497825-51-1 CAPLUS

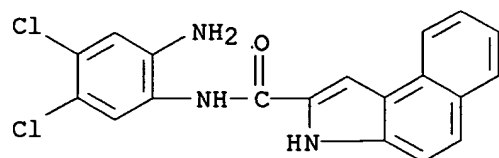
CN 1H-Indole-2-carboxamide, N-(2-amino-4,5-dichlorophenyl)-5-(2-propenyloxy)-
(9CI) (CA INDEX NAME)



RN 497825-52-2 CAPLUS

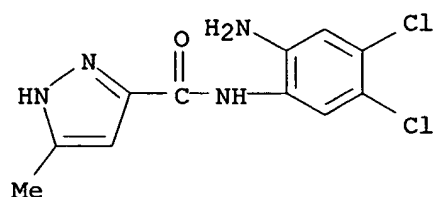
CN 3H-Benz[e]indole-2-carboxamide, N-(2-amino-4,5-dichlorophenyl)- (9CI) (CA
INDEX NAME)

10/823,211



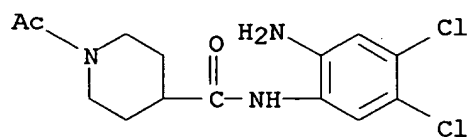
RN 497825-53-3 CAPLUS

CN 1H-Pyrazole-3-carboxamide, N-(2-amino-4,5-dichlorophenyl)-5-methyl- (9CI)
(CA INDEX NAME)



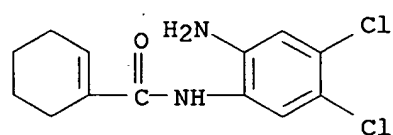
RN 497825-54-4 CAPLUS

CN 4-Piperidinecarboxamide, 1-acetyl-N-(2-amino-4,5-dichlorophenyl)- (9CI)
(CA INDEX NAME)



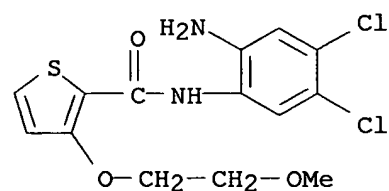
RN 497825-55-5 CAPLUS

CN 1-Cyclohexene-1-carboxamide, N-(2-amino-4,5-dichlorophenyl)- (9CI) (CA
INDEX NAME)



RN 497825-56-6 CAPLUS

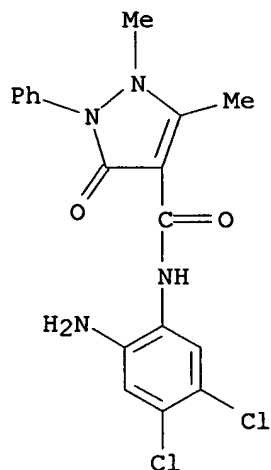
CN 2-Thiophenecarboxamide, N-(2-amino-4,5-dichlorophenyl)-3-(2-methoxyethoxy)-
(9CI) (CA INDEX NAME)



RN 497825-57-7 CAPLUS

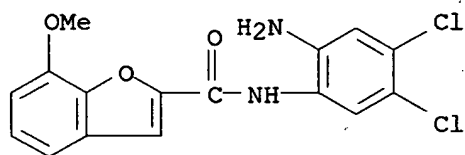
10/823,211

CN 1H-Pyrazole-4-carboxamide, N-(2-amino-4,5-dichlorophenyl)-2,3-dihydro-1,5-dimethyl-3-oxo-2-phenyl- (9CI) (CA INDEX NAME)



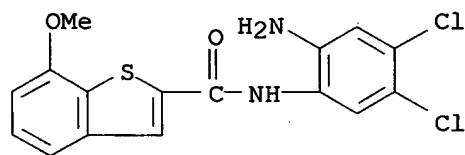
RN 497825-58-8 CAPLUS

CN 2-Benzofurancarboxamide, N-(2-amino-4,5-dichlorophenyl)-7-methoxy- (9CI)
(CA INDEX NAME)



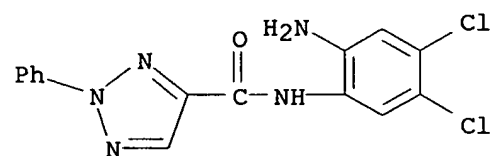
RN 497825-59-9 CAPLUS

CN Benzo[b]thiophene-2-carboxamide, N-(2-amino-4,5-dichlorophenyl)-7-methoxy- (9CI) (CA INDEX NAME)



RN 497825-60-2 CAPLUS

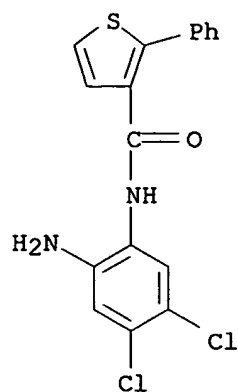
CN 2H-1,2,3-Triazole-4-carboxamide, N-(2-amino-4,5-dichlorophenyl)-2-phenyl- (9CI) (CA INDEX NAME)



RN 497825-61-3 CAPLUS

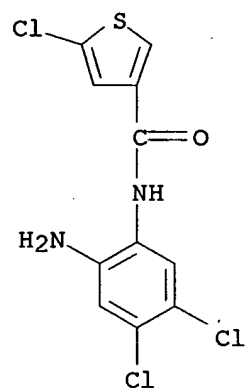
CN 3-Thiophenecarboxamide, N-(2-amino-4,5-dichlorophenyl)-2-phenyl- (9CI)
(CA INDEX NAME)

10/823,211



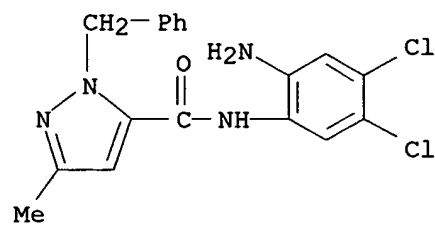
RN 497825-62-4 CAPLUS

CN 3-Thiophenecarboxamide, N-(2-amino-4,5-dichlorophenyl)-5-chloro- (9CI)
(CA INDEX NAME)



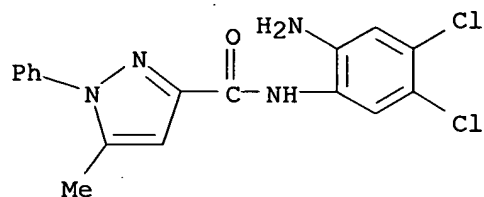
RN 497825-63-5 CAPLUS

CN 1H-Pyrazole-5-carboxamide, N-(2-amino-4,5-dichlorophenyl)-3-methyl-1-(phenylmethyl)- (9CI) (CA INDEX NAME)



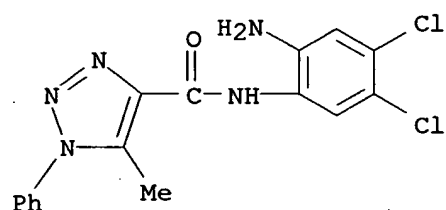
RN 497825-64-6 CAPLUS

CN 1H-Pyrazole-3-carboxamide, N-(2-amino-4,5-dichlorophenyl)-5-methyl-1-phenyl- (9CI) (CA INDEX NAME)



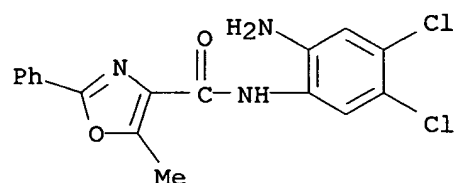
RN 497825-65-7 CAPLUS

CN 1H-1,2,3-Triazole-4-carboxamide, N-(2-amino-4,5-dichlorophenyl)-5-methyl-1-phenyl- (9CI) (CA INDEX NAME)



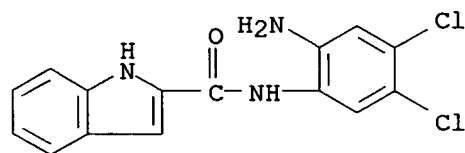
RN 497825-66-8 CAPLUS

CN 4-Oxazolecarboxamide, N-(2-amino-4,5-dichlorophenyl)-5-methyl-2-phenyl- (9CI) (CA INDEX NAME)



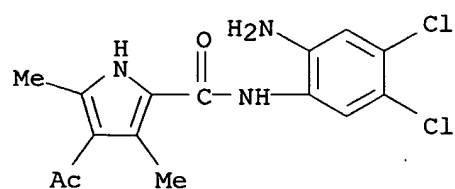
RN 497825-67-9 CAPLUS

CN 1H-Indole-2-carboxamide, N-(2-amino-4,5-dichlorophenyl)- (9CI) (CA INDEX NAME)



RN 497825-68-0 CAPLUS

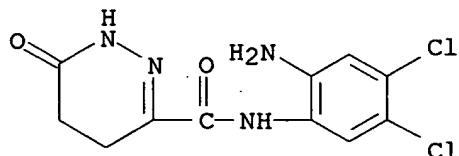
CN 1H-Pyrrole-2-carboxamide, 4-acetyl-N-(2-amino-4,5-dichlorophenyl)-3,5-dimethyl- (9CI) (CA INDEX NAME)



10/823,211

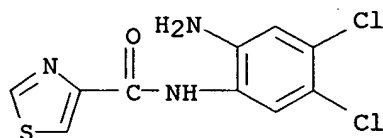
RN 497825-69-1 CAPLUS

CN 3-Pyridazinecarboxamide, N-(2-amino-4,5-dichlorophenyl)-1,4,5,6-tetrahydro-6-oxo- (9CI) (CA INDEX NAME)



RN 497825-70-4 CAPLUS

CN 4-Thiazolecarboxamide, N-(2-amino-4,5-dichlorophenyl)- (9CI) (CA INDEX NAME)



L4 ANSWER 8 OF 14 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2002:422943 CAPLUS

DN 137:6177

TI Preparation of phenylbenzimidazoles as osteoclast differentiation induction inhibitors and osteoclast inhibitors

IN Nakahira, Hiroyuki; Horiuchi, Yoshihiro

PA Sumitomo Pharmaceuticals Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 87 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	JP 2002161084	A2	20020604	JP 2000-360964	20001128
PRAI	JP 2000-360964		20001128		
OS	MARPAT 137:6177				
IT	433298-97-6 433299-06-0 433299-22-0 433299-33-3 433299-37-7				

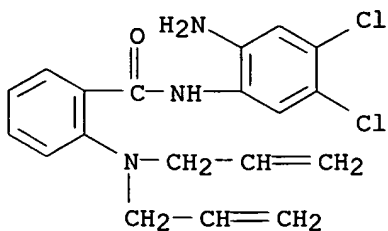
RL: RCT (Reactant); RACT (Reactant or reagent)

(preparation of phenylbenzimidazoles as osteoclast differentiation induction inhibitors and osteoclast inhibitors)

RN 433298-97-6 CAPLUS

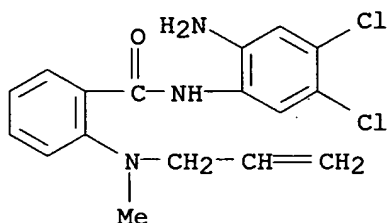
CN Benzamide, N-(2-amino-4,5-dichlorophenyl)-2-(di-2-propenylamino)- (9CI) (CA INDEX NAME)

10/823,211



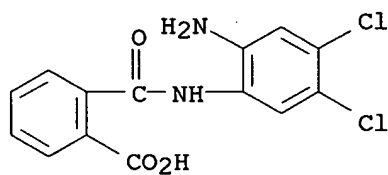
RN 433299-06-0 CAPLUS

CN Benzamide, N-(2-amino-4,5-dichlorophenyl)-2-(methyl-2-propenylamino)-
(9CI) (CA INDEX NAME)



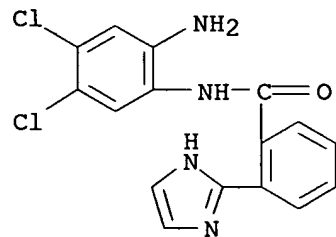
RN 433299-22-0 CAPLUS

CN Benzoic acid, 2-[[(2-amino-4,5-dichlorophenyl)amino]carbonyl]- (9CI) (CA
INDEX NAME)



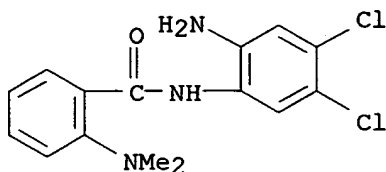
RN 433299-33-3 CAPLUS

CN Benzamide, N-(2-amino-4,5-dichlorophenyl)-2-(1H-imidazol-2-yl)- (9CI) (CA
INDEX NAME)



RN 433299-37-7 CAPLUS

CN Benzamide, N-(2-amino-4,5-dichlorophenyl)-2-(dimethylamino)- (9CI) (CA
INDEX NAME)

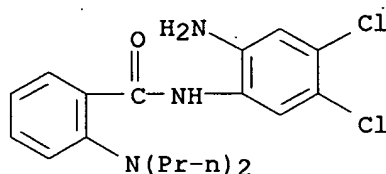
IT **433299-49-1P**

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of phenylbenzimidazoles as osteoclast differentiation induction inhibitors and osteoclast inhibitors)

RN 433299-49-1 CAPLUS

CN Benzamide, N-(2-amino-4,5-dichlorophenyl)-2-(diethylamino)- (9CI) (CA INDEX NAME)



L4 ANSWER 9 OF 14 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2001:12425 CAPLUS

DN 134:86246

TI Preparation of azolylbenzamides and analogues for treating osteoporosis

IN Farina, Carlo; Gagliardi, Stefania; Rahman, Shahzad Sharooq

PA Smithkline Beecham P.L.C., UK; Smithkline Beecham S.P.A.

SO PCT Int. Appl., 92 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001000587	A1	20010104	WO 2000-EP5881	20000623
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
	RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			
	EP 1187813	A1	20020320	EP 2000-947877	20000623
	R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO			
	JP 2003503390	T2	20030128	JP 2001-506997	20000623
	US 2005038095	A1	20050217	US 2004-868090	20040614
PRAI	GB 1999-14825	A	19990624		
	WO 2000-EP5881	W	20000623		
	US 2001-19105	A1	20011220		
OS	MARPAT 134:86246				
IT	316810-05-6P , N-(2-Amino-4,5-dichlorophenyl)-2-methoxy-4-				

10/823,211

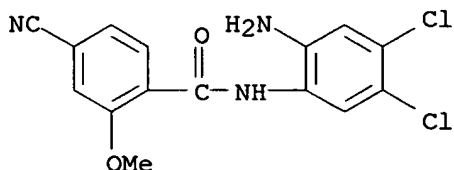
cyanobenzamide **316810-13-6P**, N-(2-Amino-4,5-dichlorophenyl)-2-ethoxy-4-cyanobenzamide

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of azolybenzamides and analogs for treating osteoporosis)

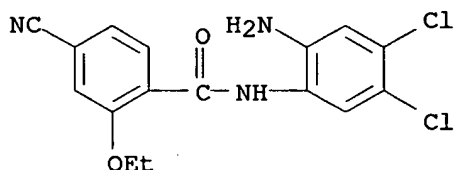
RN 316810-05-6 CAPLUS

CN Benzamide, N-(2-amino-4,5-dichlorophenyl)-4-cyano-2-methoxy- (9CI) (CA INDEX NAME)



RN 316810-13-6 CAPLUS

CN Benzamide, N-(2-amino-4,5-dichlorophenyl)-4-cyano-2-ethoxy- (9CI) (CA INDEX NAME)



RE.CNT 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 10 OF 14 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2000:68330 CAPLUS

DN 132:107785

TI Preparation of phenylenediamine compounds as macrophage scavenger receptor antagonists

IN Weinstock, Joseph; Franz, Robert G.

PA SmithKline Beecham Corporation, USA

SO PCT Int. Appl., 16 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

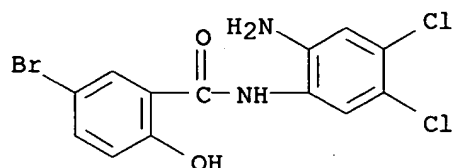
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000003704	A1	20000127	WO 1999-US16347	19990720
	W: CA, JP, US				
	RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
	CA 2338122	AA	20000127	CA 1999-2338122	19990720
	EP 1100484	A1	20010523	EP 1999-935715	19990720
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	JP 2003521446	T2	20030715	JP 2000-559839	19990720
PRAI	US 1998-93361P	P	19980720		
	WO 1999-US16347	W	19990720		
OS	MARPAT 132:107785				
IT	220340-68-1P				

10/823,211

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
(Reactant or reagent)
(preparation of phenylenediamine compds. as macrophage scavenger receptor
antagonists)

RN 220340-68-1 CAPLUS

CN Benzamide, N-(2-amino-4,5-dichlorophenyl)-5-bromo-2-hydroxy- (9CI) (CA
INDEX NAME)



RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 11 OF 14 CAPLUS COPYRIGHT 2005 ACS on STN

AN 1999:130585 CAPLUS

DN 130:168111

TI Preparation of salicylanilides as macrophage scavenger receptor
antagonists for use in treatment of cardiovascular diseases.

IN Lysko, Paul G.; Weinstock, Joseph

PA SmithKline Beecham Corporation, USA

SO PCT Int. Appl., 16 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9907382	A1	19990218	WO 1998-US16374	19980806
	W: CA, JP, US				
	RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
	CA 2298824	AA	19990218	CA 1998-2298824	19980806
	EP 1005347	A1	20000607	EP 1998-940795	19980806
	R: BE, CH, DE, ES, FR, GB, IT, LI, NL				
	JP 2001513503	T2	20010904	JP 2000-506972	19980806
	US 6255298	B1	20010703	US 2000-463863	20000128
PRAI	US 1997-55397P	P	19970806		
	WO 1998-US16374	W	19980806		

OS MARPAT 130:168111

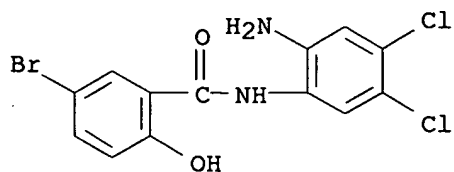
IT 220340-68-1

RL: BAC (Biological activity or effector, except adverse); BSU (Biological
study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES
(Uses)

(preparation of salicylanilides as macrophage scavenger receptor antagonists
for use in treatment of cardiovascular diseases)

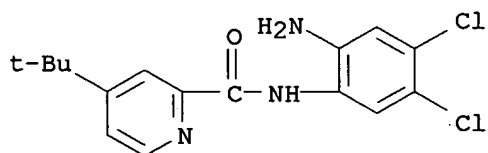
RN 220340-68-1 CAPLUS

CN Benzamide, N-(2-amino-4,5-dichlorophenyl)-5-bromo-2-hydroxy- (9CI) (CA
INDEX NAME)



RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 12 OF 14 CAPLUS COPYRIGHT 2005 ACS on STN
AN 1998:599126 CAPLUS
DN 130:32182
TI Nitrido-ruthenium(VI) and -osmium(VI) complexes containing chelating
multianionic (N,O) ligands. Synthesis, crystal structures and reactions
with triphenylphosphine
AU Chan, Pui-Ming; Yu, Wing-Yiu; Che, Chi-Ming; Cheung, Kung-Kai
CS Department of Chemistry, The University of Hong Kong, Hong Kong
SO Journal of the Chemical Society, Dalton Transactions: Inorganic Chemistry
(1998), (19), 3183-3190
CODEN: JCDBTI; ISSN: 0300-9246
PB Royal Society of Chemistry
DT Journal
LA English
IT 216394-32-0, N-(2-Amino-4,5-dichlorophenyl)-4-tert-butylpyridine-2-
carboxamide
RL: RCT (Reactant); RACT (Reactant or reagent)
(for preparation of ruthenium and osmium nitrido
pyridinecarboxamido(hydroxybenzamido)benzenato complexes)
RN 216394-32-0 CAPLUS
CN 2-Pyridinecarboxamide, N-(2-amino-4,5-dichlorophenyl)-4-(1,1-
dimethylethyl)- (9CI) (CA INDEX NAME)



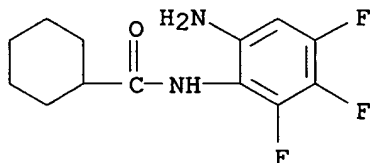
RE.CNT 40 THERE ARE 40 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 13 OF 14 CAPLUS COPYRIGHT 2005 ACS on STN
AN 1994:473886 CAPLUS
DN 121:73886
TI diaminobenzene derivatives as phospholipase A2 inhibitors, inflammation
inhibitors, and antipaneatitis agents
IN Shigehara, Itaru; Odawara, Shinji; Yuki, Shunji; Kimura, Hirohiko; Kume,
Takashi; Nakayama, Hitoshi
PA Ishihara Sangyo Kaisha, Japan
SO Jpn. Kokai Tokkyo Koho, 19 pp.
CODEN: JKXXAF
DT Patent
LA Japanese
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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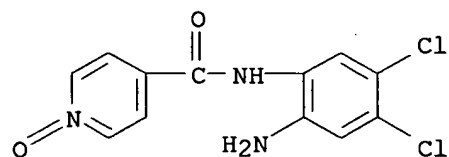
10/823,211

PI JP 06122669 A2 19940506 JP 1992-361996 19921225
PRAI JP 1991-361510 A 19911228
OS MARPAT 121:73886
IT **156522-18-8P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of, as phospholipase A2 inhibitor and inflammation inhibitor
and antipancreatitis agent)
RN 156522-18-8 CAPLUS
CN Cyclohexanecarboxamide, N-(6-amino-2,3,4-trifluorophenyl)- (9CI) (CA
INDEX NAME)



L4 ANSWER 14 OF 14 CAPLUS COPYRIGHT 2005 ACS on STN
AN 1986:608768 CAPLUS
DN 105:208768
TI N-Arylpyridine-4-carboxamides and their N-oxides
IN Crossley, Roger; Cliffe, Ian Anthony
PA John Wyeth and Brother Ltd., UK
SO Brit. UK Pat. Appl., 12 pp.
CODEN: BAXXDU
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	GB 2165537	A1	19860416	GB 1985-21304	19850827
	GB 2165537	B2	19870729		
	US 4728658	A	19880301	US 1985-771949	19850903
PRAI	GB 1984-22461	A	19840905		
OS	CASREACT 105:208768				
IT	105101-30-2P				
	RL: SPN (Synthetic preparation); PREP (Preparation)				
	(preparation of, as antisecretory and antiulcer)				
RN	105101-30-2 CAPLUS				
CN	4-Pyridinecarboxamide, N-(2-amino-4,5-dichlorophenyl)-, 1-oxide (9CI) (CA INDEX NAME)				



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'U' IS NOT VALID HERE
For an explanation, enter "HELP LOGOFF".

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COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

47.91

209.88

STN INTERNATIONAL LOGOFF AT 13:12:21 ON 09 MAY 2005